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```
10-MAY-2000
                                                   BCT
                       33801 bp
LOCUS
            Streptomyces coelicolor cosmid C77.
DEFINITION
ACCESSION
            AL136503
VERSION
            AL136503.1 GI:6714747
            adenosine deaminase; carbohydrate kinase; dehydratase;
KEYWORDS
            dihydrodipicolinate synthase; DNA-binding protein; DnaJ protein;
            dnaJ2; Era-like GTP-binding protein; GTP-binding protein;
            heat-inducible transcriptional repressor; Hit-family protein; hrcA;
            hydrolase; IclR-family transcriptional regulator; lepA;
            lipoprotein; long-chain fatty-acid CoA ligase; oxidoreductase;
            oxygen-independent coproporphyrinogen III oxidase; protease;
            transmembrane efflux protein; transmembrane transport protein.
            Streptomyces coelicolor A3(2).
SOURCE
            Streptomyces coelicolor A3(2)
  ORGANISM
            Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
            Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.
               (bases 1 to 33801)
REFERENCE
            Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J.,
  AUTHORS
            Kinashi, H. and Hopwood, D.A.
            A set of ordered cosmids and a detailed genetic and physical map
  TITLE
            for the 8 Mb Streptomyces coelicolor A3(2) chromosome
           Mol. Microbiol. 21 (1), 77-96 (1996)
  JOURNAL
            97000351
  MEDLINE
REFERENCE
               (bases 1 to 33801)
           Oliver, K. and Harris, D.
  AUTHORS
  JOURNAL
           Unpublished
               (bases 1 to 33801)
REFERENCE
           Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A.
  AUTHORS
           Direct Submission
  TITLE
  JOURNAL
           Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project,
           Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge
           CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof.
            David A. Hopwood, [3] John Innes Centre, Norwich Research Park,
           Colney, Norwich, Norfolk NR4 7UH, UK
alignment scores:
                                        Length:
                                                   363
           Ouality:
                     397.00
                                          Gaps:
              Ratio:
                     1.829
                              Percent Identity: 33.609
                     59.780
Percent Similarity:
alignment block:
US-09-477-962-115 \times SCC77/rev
Align seg 1/1 to reverse of: SCC77 from: 1 to: 33801
      11 IleProAlaIleArgGluAlaLeuGlyAspGluLysAspProArgLeuAl 27
                       :::||||||
                                           111
         :::|||||
   26678 CTGCCCGCG.....TCCGCGCTCGCCGGGGCCGCCGACCGCCCCCTCGG 26635
      27 aLeuTyrValHisValProPheCysSerSerLysCysHisPheCysAspT 44
         694 GTTCTACCTGCACGTGCCGTACTGCGCGACCCGCTGCGGCTACTGCGACT 26585
         rpValThrAspIleProValAlaArgLeuArgGlyAspSer
```

26584 TGAACAGC TACACCGCCACCGAACTGCGCGGCACCGGCGGGGGTCCTC 26538

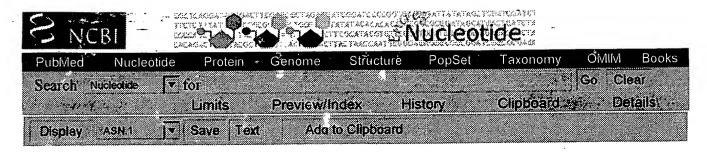
A111

:::

All Control of the Co

	ArgGluArgSerProTyrValThrAlaLeuCysAspGlnIleArgPheTy ::: ::: ::: ::: :::::	
	GCCTCCCGCGACAACTACGCCGACACCCTCGTCGACGAGGTCCGC	
	rGlyProGlnLeuThrArgLeuGlyTyrArgProG	
•	CTGGCCCGCAAGGTGCTCGGCGACGACCCCCGCGAGGTCC	
	<pre>luValMetTyrTrpGlyGlyGlyThrProThrArgLeuThrGlyAspGlu ::::::::::::::::::::::::::::::::::::</pre>	
26452	GCACGGTCTTCGTCGGCGGCGGTACGCCCACCCTGCTGGCCGCCGGCGAC	26403
	MetThrAlaValHisGlnAlaLeuAspAspAlaPheAspLeuThrGlyLe ::::: :: ::: :::	
26402	CTGGTGCGGATGCTGGGCGCGATCCGCGACGAGTTCGGCCTGGCACCGGA	26353
	"uArgGlnTrpSerValGluSerThrProAsnAspLeuAspProAlaThrL	:
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136	<pre>euAspThrLeuArgGlyLeuGlyValThrArgValSerValGlyValGln ::: ::: ::: </pre>	152
26302	TCGCCACCTCCGCGCGGGCGGCTTCAACCGGATCTCCTTCGGCATGCAG	26253
153	SerLeuAsnProTyrGlnLeuArgLysAlaGlyArgAlaHisSerArgGl	169
26252	AGCGCCAAGCACGCCCTGAAGATCCTGGACCGCACCCCACACCCCGGG	26203
169	uGlnAlaLeuAlaAlaValProLeuLeuArgArgAlaGlyIleAspGluP ::: ::: ::: :::	186
26202	ACGCCCCGAGGCCTGTGTCGCCGAGGCCCGCGCGGCCGGC	26153
186	heAsnValAspLeuIleAlaGlyPheProGlyGluAlaValGluSerPhe	202
26152	TCAACCTCGACCTGATCTACGGCACCCCCGGCGAGTCCGACGACGACTGG	26103
203	GluGluThrLeuArgThrValLeuAlaLeuAspProProHisValSerVa ::: ::: ::::: ::: :	219
26102	CGGGCCTCCCTGGACGCCGCGCCCGGACCCGACCACGTCTCGGC	26053
219	<pre>lTyrProTyrArgAlaThrProLysThrValMetAlaMetGlnLeuAspA : </pre>	236
26052	GTACGCCCTGATCGTCGAGGAGGGCACCCAGCTCGCCCGCC	26003
236	rgGluPheValGluAlaArgAsnArgAspGlyMetIleAspAlaTyrGlu	252
26002	GCGGCGAGGTCCCGATGACCGACGACGACGTGCACGCCGACCGGTACCTG	25953
253	ArgAlaMetAlaAlaLeuGlyAlaAlaGlyTyrHisGluTyrCysHisGl	269
	ATCGCCGAGGAGGCACTGTCCGCGGCCGGTTACGACTGGTACGAGGTGTC	
269	yTyrTrpValArgAspAlaArgHisGluAspGlnAspG	282
	CAACTGGGCCACCTCCGACGCGGGGCGCTGCCTGCAC	25866
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282	<pre>lyAsnTyrLysTyrAspLeuAlaGlyAspLysIleGlyPheGlySerGly </pre>	298
25865	AACGAGCTGTACTGGCGGGGCGCCGACTGGTGGGGCCCGGACCGGGC	25818
299	AlaGluSerIleIleGlyHisHisLeuLeuTrpAsnGluAsnSe	313
25817	GCGCACTCCCACGTGGGGGGCGTGCGGTGGTGGAACGTGAAGCACCCGGG	25768
313	rAlaTyrAlaArgTyrLeuLeuAlaProArgGluPheSerAlaAlaHisA: :::::::::::::::::::::::::	330
25767	GGCGTACGCGGGGGCGCGGGCGCGGGCGCGGGCGCG	25718
330	rgPheThrThrAlaGluProAspArgLeuThrAlaProValGlyGlyAla ::::: ::: :::	346
25717	AGATCCTCACGGACGAGGACCGGCGCGTGGAGCGCATCCTGCTGGAG	25671
347	LeuMetThrArgGluGlyValValPheAlaArgPheArg 359	
25670	CTGCGCCTGCGGGAGGGCGTCCCGCTGTCGCTGCTGCGG 25632	



Related Sequences, Protein, PubMed, Taxonomy **1:** AL136503. Streptomyces coel...[gi:6714747] BCT 10-MAY-2000 linear 33801 bp DNA LOCUS Streptomyces coelicolor cosmid C77. DEFINITION AL136503 ACCESSION **VERSION** AL136503.1 GI:6714747 adenosine deaminase; carbohydrate kinase; dehydratase; **KEYWORDS** dihydrodipicolinate synthase; DNA-binding protein; DnaJ protein; dnaJ2; Era-like GTP-binding protein; GTP-binding protein; heat-inducible transcriptional repressor; Hit-family protein; hrcA; hydrolase; IclR-family transcriptional regulator; lepA; lipoprotein; long-chain fatty-acid CoA ligase; oxidoreductase; oxygen-independent coproporphyrinogen III oxidase; protease; transmembrane efflux protein; transmembrane transport protein. Streptomyces coelicolor A3(2). SOURCE Streptomyces coelicolor A3(2) ORGANISM Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces. REFERENCE (bases 1 to 33801) Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J., AUTHORS Kinashi, H. and Hopwood, D.A. A set of ordered cosmids and a detailed genetic and physical map TITLE for the 8 Mb Streptomyces coelicolor A3(2) chromosome Mol. Microbiol. 21 (1), 77-96 (1996) **JOURNAL** MEDLINE 97000351 REFERENCE (bases 1 to 33801) Oliver, K. and Harris, D. AUTHORS JOURNAL Unpublished REFERENCE (bases 1 to 33801) Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A. AUTHORS TITLE . Direct Submission Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project, **JOURNAL** Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof. David A. Hopwood, [3] John Innes Centre, Norwich Research Park, Colney, Norwich, Norfolk NR4 7UH, UK COMMENT Notes: Streptomyces coelicolor sequencing at The Sanger Centre is funded by the BBSRC and Beowulf Genomics Details of S. coelicolor sequencing at the Sanger Centre are available on the World Wide Web. (URL; http://www.sanger.ac.uk/Projects/S_coelicolor/) CDS are numbered using the following system eg SC7B7.01c. SC (S. coelicolor), 7B7 (cosmid name), .01 (first CDS), c (complementary strand). The more significant matches with motifs in the PROSITE database are also included but some of these may be fortuitous. The length in codons is given for each CDS. Usually the highest scoring match found by fasta -o is given for CDS which show significant similarity to other CDS in the database.

The position of possible ribosome binding site sequences are given

Gene prediction is based on positional base preference in codons

where these have been used to deduce the initiation codon.

S NO	TBI Nucleo		Genome	Structure	PopSet	tein Taxonomy	OMIM	Books
	Nucleatide	for Limits	Preview/Ind		istory	Clipboard	Go Cle	er tails
Display	default	▼ Save T	ext Add to	Clipboard				
☐1: CAB66237. putative oxygen-i[gi:6714773] Nucleotide, Related Sequences, PubMed, Taxonomy BLink, LinkOut								

BCT 10-MAY-2000 LOCUS CAB66237 435 aa linear DEFINITION putative oxygen-independent coproporphyrinogen III oxidase. [Streptomyces coelicolor A3(2)]. ACCESSION CAB66237 PID g6714773 VERSION CAB66237.1 GI:6714773 **DBSOURCE** embl locus SCC77, accession AL136503.1 **KEYWORDS** Streptomyces coelicolor A3(2). SOURCE Streptomyces coelicolor A3(2) ORGANISM Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces. REFERENCE 1 (residues 1 to 435) Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J., AUTHORS Kinashi, H. and Hopwood, D.A. A set of ordered cosmids and a detailed genetic and physical map TITLE for the 8 Mb Streptomyces coelicolor A3(2) chromosome JOURNAL Mol. Microbiol. 21 (1), 77-96 (1996) MEDITNE 97000351 REFERENCE (residues 1 to 435) AUTHORS Oliver, K. and Harris, D. JOURNAL Unpublished REFERENCE (residues 1 to 435) Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A. **AUTHORS** TITLE Direct Submission JOURNAL Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project, Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof. David A. Hopwood, [3] John Innes Centre, Norwich Research Park, Colney, Norwich, Norfolk NR4 7UH, UK COMMENT Notes: Streptomyces coelicolor sequencing at The Sanger Centre is funded by the BBSRC and Beowulf Genomics Details of S. coelicolor sequencing at the Sanger Centre are available on the World Wide Web. (URL; http://www.sanger.ac.uk/Projects/S_coelicolor/) CDS are numbered using the following system eg SC7B7.01c. SC (S. coelicolor), 7B7 (cosmid name), .01 (first CDS), c (complementary strand). The more significant matches with motifs in the PROSITE database are also included but some of these may be fortuitous. The length in codons is given for each CDS. Usually, the highest scoring match found by fasta -o is given for. CDS which show significant similarity to other CDS in the database. The position of possible ribosome binding site sequences are given where these have been used to deduce the initiation codon. Gene prediction is based on positional base preference in codons using a specially developed Hidden Markov Model (Krogh et al., Nucleic Acids Research, 22(22):4768-4778(1994)) and the FramePlot program of Bibb et al., Gene 30:157-66(1984) as implemented at

1 of 2

```
http://www.nih.go.jp/
            jun/cgi-bin/frameplot.pl. CAUTION: We may not have predicted the
            correct initiation codon. Where possible we choose an initiation
            codon (atg, gtg, ttg or (att)) which is preceded by an upstream
            ribosome binding site sequence (optimally 5-13bp before the
            initiation codon). If this cannot be identified we choose the most
            upstream initiation codon.
            IMPORTANT: This sequence MAY NOT be the entire insert of the
            sequenced clone. It may be shorter because we only sequence
            overlapping sections once, or longer, because we arrange for a
            small overlap between neighbouring submissions.
            Cosmid C77 Lies between and overlaps with cosmids C117 and C123 on
            the AseI-C genomic restriction fragment.
FEATURES
                     Location/Qualifiers
                     1..435
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                     /organism="Streptomyces coelicolor A3(2)"
                     /strain="A3(2)"
                     /db xref="taxon:100226"
                     /clone="cosmid C77"
     Protein
                     1..435
                     /product="putative oxygen-independent coproporphyrinogen
                     III oxidase."
     CDS
                     1..435
                     /gene="SCC77.26c"
                     /coded by="complement(AL136503.1:25494..26801)"
                     /transl table=11
                     /note="SCC77.26c, possible oxygen-independent
                     coproporphyrinogen III oxidase (EC 1.-.-.), len: 435 aa.
                     Highly similar to many putative coproporphyrinogen III
                     oxidases including: Bacillus subtilis
                     SW:HEMN BACSU(EMBL:X91655) probable oxygen-independent
                     coproporphyrinogen III oxidase (366 aa), fasta scores opt:
                     490 z-score: 555.3 E(): 1.5e-23 30.8% identity in 328 aa
                     overlap and Mycobacterium tuberculosis
                     SW:HEMN MYCTU(EMBL:Z81368) probable oxygen-independent
                     coproporphyrinogen III oxidase (375 aa), fasta scores opt:
                     1358 z-score: 1530.5 E():0 56.5% identity in 382 aa
                     overlap. Contains a Prosite hit to PS00017 ATP/GTP-binding
                     site motif A (P-loop)."
ORIGIN
        1 mngrreraqg tewagdpagc gtmermpsal pdgepvpadg alpasalaga adrplgfylh
       61 vpycatrcgy cdfntytate lrgtggvlas rdnyadtlvd evrlarkvlg ddprevrtvf
      121 vgggtptlla agdlvrmlga irdefglapd aeitteanpe svdpaylatl raggfnrisf
      181 gmqsakqhvl kildrthtpg rpeacvaear aagfdhvnld liygtpgesd ddwrasldaa
     241 lgagpdhvsa yaliveegtq larrirrgev pmtdddvhad ryliaeeals aagydwyevs
     301 nwatsdagrc lhnelywrga dwwgagpgah shvggvrwwn vkhpgayaga laagkspgag
     361 reiltdedrr verillelrl regvplsllr eaglaasrra lsegllqegp yeagsavltl
      421 rgrlladavv rdlvd
11
```

Revised: October 24, 2001.

5 x US-09-477-962-1 ...

Seg 1= 58857 6p 57,583-58,854

alignment block: $us-09-477-962-115 \times US-09-477-962-1$ Align seg 1/1 to: US-09-477-962-1 from: 1 to: 58857 1 MetSerHisAlaIleGlyProSerArgLeuIleProAlaIleArgGluAl 17 57583 ATGAGCCACGCCATCGGAPCGAGCCGGCTGATCCCCGCCATCCGCGAAGC 57632 17 aLeuGlyAspGluLysAspProArgLeuAlaLeuTyrValHisValProP 34 57633 GCTCGGGGACGAGAAGGACCCCCGGCTCGCCCTCTACGTCCACGTCCCCT 57682 34 heCysSerSerLysCysHisPheCysAspTrpValThrAspIleProVal 50 57683 TCTGCTCCTCCAAGTGCCACTTCTGCGACTGGGTCACCGACATCCCCGTC 57732 51 AlaArgLeuArgGlyAspSerArgGluArgSerProTyrValThrAlaLe 67 57733 GCACGCCTGCGCGGCACAGCCGGGAACGCTCGCCCTACGTCACCGCCCT 57782 67 uCysAspGlnIleArgPheTyrGlyProGlnLeuThrArgLeuGlyTyrA 84 57783 CTGCGACCAGATCCGCTTCTACGGCCCCCAGCTCACCCGGCTCGGCTACC 57832 84 rgProGluValMetTyrTrpGlyGlyGlyThrProThrArgLeuThrGly 100 101 AspGluMetThrAlaValHisGlnAlaLeuAspAspAlaPheAspLeuTh 117 57883 GACGAGATGACGGCCGTCCACCAGGCCCTCGACGACGCCTTCGACCTGAC 57932 117 rGlyLeuArgGlnTrpSerValGluSerThrProAsnAspLeuAspProA 134 57933 GGGACTCCGCCAGTGGTCGGTGGAGAGCACCCCGAACGACCTCGACCCCG 57982 134 laThrLeuAspThrLeuArgGlyLeuGlyValThrArgValSerValGly 150 57983 CCACCCTCGACACCCTGCGCGCGCCTCGGCGTCACCCGCGTCAGCGTCGGC 58032 151 ValGlnSerLeuAsnProTyrGlnLeuArgLysAlaGlyArgAlaHisSe 167 58033 GTCCAGTCGCTCAACCCGTACCAGCTGCGCAAGGCAGGCCGGGCCCACTC 58082 167 rArgGluGlnAlaLeuAlaAlaValProLeuLeuArgArgAlaGlyIleA 184 58083 GCGCGAACAGGCCCTGGCCGCCGTCCCCCTGTTGCGCCGCGCCGCGCATCG 58132 184_spGluPheAsnValAspLeuIleAlaGlyPheProGlyGluAlaValGlu 200 cactificaacctecaccticaticgccccctificgccggccaacccticgac.

201 SerPheGluGluThrLeuArgThrValLeuAlaLeuAspProProHisVa 217

217	lSerValTyrProTyrArgAlaThrProLysThrValMetAlaMetGlnL	234
- -58233	CTCCGTCTACCCCTACCGCGCCACCCCCAAGACGGTCATGGCCATGCAGC	58282
234	euAspArgGluPheValGluAlaArgAsnArgAspGlyMetIleAspAla	250
58283	TCGACCGCGAGTTCGTCGAGGCCCGGAACCGGGACGGCATGATCGACGCC	58332
251	TyrGluArgAlaMetAlaAlaLeuGlyAlaAlaGlyTyrHisGluTyrCy	267
58333	TATGAACGGCCCATGGCCGCCGCCTCGCCGCCGCCTATCACGAGTACTG	58382
267	sHisGlyTyrTrpValArgAspAlaArgHisGluAspGlnAspGlyAsnT	284
58383	CCACGGCTACTGGGTGCGCGACGCGCCACGAGGACCAGGACGCCAACT	58432
284	<pre>yrLysTyrAspLeuAlaGlyAspLysIleGlyPheGlySerGlyAlaGlu</pre>	300
58433	ACAAGTACGACCTGGCCGGCGACAAGATCGGCTTTGGCAGCGGCGCCGAA	58482
301	SerIleIleGlyHisHisLeuLeuTrpAsnGluAsnSerAlaTyrAlaAr	317
58483	TCGATCATCGGTCACCACCTGCTCTGGAACGAGAACAGCGCCTACGCCCG	58532
317	gTyrLeuLeuAlaProArgGluPheSerAlaAlaHisArgPheThrThrA	334
58533	CTACCTGCTCGCCCCGCGAGTTCTCCGCCGCCCACCGGTTCACCACCG	58582
334	laGluProAspArgLeuThrAlaProValGlyGlyAlaLeuMetThrArg	350
58583	CCGAACCGACCGCCTGACCGCCCCCGTCGGCGCGCGCTGATGACCCGT	58632
351	GluGlyValValPheAlaArgPheArgArgLeuThrGlyLeuAspPheAl	3.67
58633	GAAGGCGTGGTCTTCGCCCGCTTCCGCAGACTGACCGGCCTGGACTTCGC	58682
367	aAspValArgAlaThrProTyrPheArgGlnTrpPheGluLeuLeuGluA	384
58683	GGACGTCCGCCACACCGTACTTCCGCCAGTGGTTCGAGCTCCTGGAGC	58732
384	rgCysGlyGlyArgPheValGluThrProTyrSerLeuArgLeuGluPro	400
58733	GCTGCGGCGGCCGCTTCGTCGAGACGCCGTACAGCCTCCGCCTGGAGCCG	58782
401	SerThrlleHisArgAlaTyrlleThrHisLeuAlaTyrThrMetAlaHi	417
58783	TCCACCATCCACCGCGCCTACATCACCCACCTCGCCTACACCATGGCCCA	58832
417	sGlyLeuAlaProGluArgAla 424	
58833	TGGCCTGGCCCCGAACGCGCC 58854	

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- L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AN 2000:332950 CAPLUS
- TI Bleomycin biosynthesis in Streptomyces verticillus ATCC15003: A model for hybrid peptide and polyketide biosynthesis.
- AU Du, Liangcheng; Sanchez, Cesar; Chen, Mei; Edwards, Daniel J.; Murrell, Jeffrey M.; Shen, Ben
- CS Department of Chemistry, University of California, Davis, CA, 95616, USA
- SO Book of Abstracts, 219th ACS National Meeting, San Francisco, CA, March 26-30, 2000 (2000), ORGN-822 Publisher: American Chemical Society, Washington, D. C.
 - CODEN: 69CLAC
- DT Conference; Meeting Abstract
- LA English

=> d abs

- L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- Polyketides and nonribosomal peptides are assembled in a remarkably similar fashion by polyketide synthases (PKSs) from short carboxylic acids and nonribosomal peptide synthetases (NRPSs) from amino acids, resp. Cloning and sequence anal. of the 90-kb bleomycin (BLM) biosynthesis cluster from Streptomyces verticillus ATCC15003 revealed both NRPS and PKS genes. By detg. the substrate specificity of individual NRPS and PKS modules, a linear hybrid NRPS/PKS/NRPS model is formulated for the Blm megasynthetase-templated assembly of BLM from nine amino acids and one acetate. These results set the stage for engineering novel BLM analogs by genetic manipulation of the blm biosynthesis genes, support the wisdom of combining individual NRPS and PKS modules for combinatorial biosynthesis, and lay the foundation to investigate the mol. basis for intermodular communication between NRPS and PKS and the mechanism for bithiazole biosynthesis.

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- L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AN 1999:145586 CAPLUS
- TI Bleomycin biosynthesis in streptomyces verticillus ATCC15003: The search for a hybrid polyketide and peptide biosynthetic system
- AU Shen, Ben; Du, Liangcheng; Edwards, Dan; Chen, Mei; Sanchez, Cesar
- CS Department of Chemistry, University of California, Davis, CA, 95616, USA
- SO Book of Abstracts, 217th ACS National Meeting, Anaheim, Calif., March 21-25 (1999), ORGN-153 Publisher: American Chemical Society, Washington, D. C.
 - CODEN: 67GHA6
- DT Conference; Meeting Abstract
- LA English

=> d abs

- L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AB The biosynthesis of bleomycin (Blm) in Sv. ATCC15003 has been studied as a model to decipher the mechanism of how peptide synthase (PTS) and polyketide synthase (PKS) can be hybridized into a functional system to make metabolite from amino acids and short fatty acids. A 110kb gene cluster for Blm biosynthesis was cloned from Sv. ATCC15003, 75kb of which has been fully sequenced and analyzed. Among the many novel discoveries made from this study are:. (1) the first model for a hybrid PTS/PKS/PTS biosynthetic system, (2) the first example of PKS gene from actinomycetes that contains a MT domain, and (3) a novel mechanism for bithiazole biosynthesis. These results should lay the foundation for rational engineering of hybrid PTS/PKS biosynthetic systems from other peptide and polyketide biosynthetic pathways to generate structural diversity.

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